

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A method for ~~extracting antineoplastic components obtaining~~
an extract from *Bupleurum scorzonerifolium*, wherein the extract comprises antineoplastic
components, the method comprising the steps of:
extracting a lignan mixture from *Bupleurum scorzonerifolium* wherein the lignan mixture
further comprises at least one antineoplastic component; and
isolating a pure *Bupleurum scorzonerifolium* extract from the lignan mixture
dissolving *Bupleurum scorzonerifolium* powder using an acetone solution to obtain an
acetone extract and residues;
dissolving the acetone extract in a protic solvent solution and extracting the protic solvent
solution with a nonpolar organic solvent, to obtain a nonpolar organic solvent extract and a protic
solvent portion; and
dissolving the protic solvent portion in an aqueous solvent and extracting with a polar
organic solvent to obtain a polar organic solvent extract and an aqueous portion.
2. (Canceled)
3. (Currently amended) The method of claim [[2]] 1, wherein the ~~first solution is~~
acetoneprotic solvent solution is 95% methanol/5% water.
4. (Currently amended) The method of claim [[2]] 1, wherein the ~~second solution is~~
methanolaqueous solvent is water.
5. (Currently amended) The method of claim [[2]] 1, wherein the ~~third solution is 95%~~
methanol solutionnonpolar organic solvent is hexane.

6. (Currently amended) The method of claim ~~[[2]]~~ 1, wherein the ~~fourth solution is chloroform~~ polar organic solvent is chloroform.

7. (Currently amended) The method of claim ~~[[2]]~~ 1, further ~~comprises~~ comprising the step of separating the antineoplastic components from the acetone extract, the nonpolar organic solvent extract, and the polar organic solvent extract by a chromatographic method for separating antineoplastic components.

8. (Original) The method of claim 7, wherein the chromatographic method is a silica gel chromatography

9. (Original) The method of claim 7, wherein the chromatographic method is a high performance liquid chromatography (HPLC).

10. (Original) The method of claim 7, wherein the chromatographic method is a medium pressure liquid chromatography (MPLC).

11. (Canceled)

12. (Currently amended) The method of claim 1, wherein ~~the cell proliferative disorder includes~~ the antineoplastic components have anti-tumor effects on hepatoma.

13. (Currently amended) The method of claim 1, wherein the antineoplastic components have anti-tumor effects on ~~cell proliferative disorder includes~~ ovarian cancer.

14. (Currently amended) The method of claim 1, wherein the antineoplastic components have anti-tumor effects on ~~cell proliferative disorder includes~~ malignant glioblastoma.

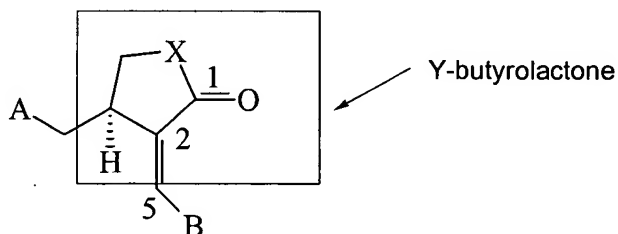
15. (Currently amended) The method of claim 1, wherein the antineoplastic components have anti-tumor effects on ~~cell proliferative disorder includes~~ lung cancer.

16. (Currently amended) The method of claim 1, wherein the antineoplastic components have anti-tumor effects on cell proliferative disorder ~~includes colorectal cancer~~.

17. (Currently amended) The method of claim 1, wherein the antineoplastic components have anti-tumor effects on cell proliferative disorder ~~is~~ tumors which are resistant to a taxane-type anticancer agent.

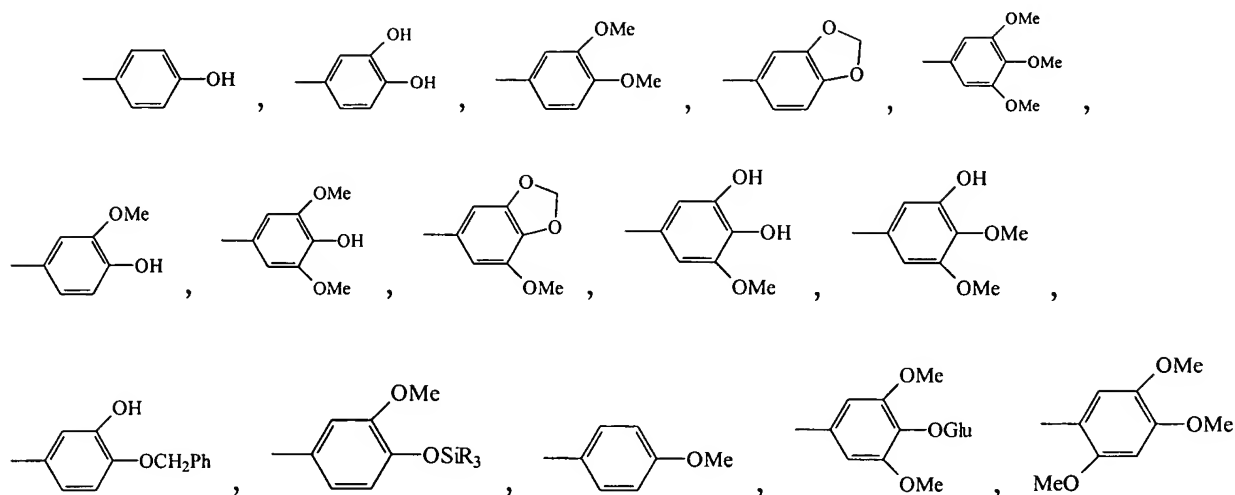
18. (Original) The method of claim 17, wherein the taxane-type anticancer agent is Paclitaxel.

19. (Withdrawn) The method of claim 1, wherein the antineoplastic components comprise at least one of the following heterocyclic compounds or pharmacologically compatible salts, esters, ketones or derivatives based on the following formula:



wherein X includes N, O, S, and Se.

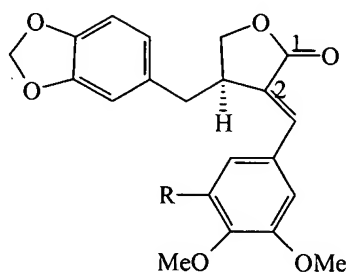
20. (Withdrawn) The method of claim 19, wherein A, B substituents can be selected from the following substituent structures:



21. (Withdrawn) The method of claim 19, wherein the heterocyclic compounds have a Z configuration at a carbon 2(5) position.

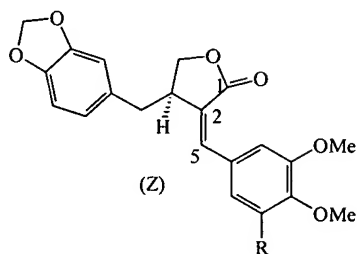
22. (Withdrawn) The method of claim 19, wherein the heterocyclic compounds have a E configuration at a carbon 2(5) position.

23. (Withdrawn) The method of claim 19, wherein the heterocyclic compound having the following formula is named chaihulactone:



where R represents a methoxyl group.

24. (Currently amended) The method of claim 1, wherein the antineoplastic components comprise a ~~the heterocyclic compound having the following formula; is named isochaihulactone:~~



where R represents a hydrogen atom, a methoxyl group, or an aromatic group.

25. (Currently amended) The method of claim 1, wherein the antineoplastic components is an agent for arresting a G2/M stage of a cell cycle.

26. (Currently amended) The method of claim 1, wherein the antineoplastic components stabilize ~~[[is]]~~ a microtubule-stabilizing agent.

27. (Currently amended) The method of claim 1, wherein the antineoplastic components are ~~[[is]]~~ used in conjunction with an anti-tumor drug.

28. (Currently amended) The method of claim 27, wherein the anti-tumor drug includes ~~is~~ Cisplatin.

29. (Currently amended) The method of claim 27, wherein the anti-tumor drug includes ~~is~~ Taxol.

30. (Currently amended) The method of claim 1, wherein the antineoplastic components is an agent for down-regulating ~~regulate~~ human telomerase reverse transcriptase (hTERT) gene.

31. (Currently amended) The method of claim 30, wherein the antineoplastic components down regulate ~~regulates~~ hTERT gene in human immunodeficiency virus (HIV).

32. (Currently amended) The method of claim 1, wherein the antineoplastic components is an agent for inhibiting telomerase activity.

33. (Currently amended) The method of claim 32, wherein the antineoplastic components ~~inhibits~~ inhibit telomerase activity in ~~the~~ HIV.